AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Original) A compound of formula (I):

and the pharmaceutically acceptable salts, and other pharmaceutically acceptable biohydrolyzable derivatives thereof;

wherein R^1 is an optionally substituted C_{3-12} carbocyclyl or C_{3-12} heterocyclyl group or group of formula (II)

wherein X is NR^3 , O, S or $(CR^{22}R^{22})_n$, Y is absent or is NR^{23} , O, or $(CR^{23}R^{23})_n$, R^2 is optionally substituted C_{1-12} alkyl, C_{2-12} alkenyl, C_{2-12} alkynyl, C_{3-12} carbocyclyl or C_{3-12} heterocyclyl, and R^4 is an optionally substituted five or six membered heterocyclyl group or an optionally substituted six membered carbocyclyl group.

2. (Original) A compound as claimed in claim 1 wherein the optionally substituted carbocyclyl or heterocyclyl group of R¹ is optionally fused to a partially saturated, unsaturated or fully saturated five to seven membered ring containing zero to three heteroatoms, and each substitutable carbon atom in R¹, including the optional fused ring, is optionally and independently substituted by one or more of halogen, C₁₋₁₂ alkyl, C₂₋₁₂ alkenyl, C₂₋₁₂ alkynyl, haloC₁₋₁₂alkyl, C₃₋₁₂ carbocyclyl, C₃₋₁₂ heterocyclyl, (CH₂)_nOR⁵, (CH₂)_nNR⁵₂(CH₂)_nSR⁵, OR⁵, SR⁵, NO₂, CN, NR⁵₂, NR⁵COR⁵, NR⁵CONR⁵₂, NR⁵COR⁵, NR⁵CO₂R⁵, CO₂R⁵, COR⁵, CONR⁵₂, S(O)₂R⁵, SONR⁵₂, S(O)₈R⁵, SO₂NR⁵₂, or NR⁵S(O)₂R⁵ wherein the C₁₋₁₂ alkyl group optionally

contains one or more insertions selected from -O-, -N(R⁵)- -S-, -S(O)- and -S(O₂)-; and each saturated carbon in the optional fused ring is further optionally and independently substituted by =O, =S, NNR⁶₂, =N-OR⁶, =NNR⁶COR⁶, =NNR⁶CO₂R⁶, =NNSO₂R⁶, or =NR⁶; and each substitutable nitrogen atom in R¹ is optionally substituted by R⁷, COR⁷, SO₂R⁷ or CO₂R⁷;

wherein n is 1 to 6, preferably n is 1, 2 or 3;

wherein R^5 is hydrogen, C_{1-12} alkyl, C_{3-12} carbocyclyl or C_{3-12} heterocyclyl, optionally substituted by one or more of C_{1-6} alkyl, C_{3-12} carbocyclyl, C_{3-12} heterocyclyl, halogen, C_{1-6} haloalkyl, OR^8 , SR^8 , NO_2 , CN, NR^8R^8 , NR^8COR^8 , NR^8COR^8 , NR^8COR^8 , $NR^8CO_2R^8$, CO_2R^8 ,

wherein two R⁵ in NR⁵₂ may optionally form a partially saturated, unsaturated or fully saturated three to seven membered ring containing one to three heteroatoms, optionally and independently substituted by one or more of C₁₋₆ alkyl, halogen, C₁₋₆ haloalkyl, OR⁸, SR⁸, NO₂, CN, NR⁸R⁸, NR⁸COR⁸, NR⁸COR⁸, NR⁸COR⁸, NR⁸CO₂R⁸, CO₂R⁸, COR⁸, CONR⁸₂, S(O)₂R⁸, SO₂NR⁸R⁸, NR⁸S(O)₂R⁸,

wherein the C_{1-6} alkyl group optionally incorporates one or two insertions selected from the group consisting of -O-, -N(R^8)-, -S(O)- and -S(O₂)-, wherein each R^8 may be the same or different and is as defined below;

wherein R^6 is hydrogen, C_{1-12} alkyl, C_{3-12} carbocyclyl or C_{3-12} heterocyclyl, optionally substituted by one or more of C_{1-6} alkyl, halogen, C_{1-6} haloalkyl, OR^8 , SR^8 , NO_2 , CN, NR^8R^8 , NR^8COR^8 , NR^8COR^8 , NR^8COR^8 , $NR^8CO_2R^8$, CO_2R^8 , CO_2R^8 , $CONR^8_2$, $S(O)_2R^8$, $S(O)_2R^8$, $S(O)_2R^8$, $S(O)_2R^8$, wherein the C_{1-12} alkyl group optionally incorporates one or two insertions selected from the group consisting of -O-, -N(R^8)-, -S(O)- and -S(O₂)-, wherein each R^8 may be the same or different and is as defined below;

wherein R⁷ is hydrogen, C₆₋₁₂ aryl, C₁₋₆ alkyl or C₁₋₆ haloalkyl;

wherein R⁸ is hydrogen, C₁₋₆ alkyl, or C₁₋₆ haloalkyl.

3. (Currently Amended) A compound as claimed in claim 1 or 2 wherein Y is absent or is NR^{23} , O, $(CR^{23}R^{23})_n$,

wherein each R²³ is H, C₁₋₄ alkyl, C₁₋₄ alkoxy or C₁₋₄ haloalkyl;

and n is 1 to 6, preferably n is 1, 2, 3 or 4; and

 R^2 is optionally substituted C_{1-12} alkyl, C_{2-12} alkenyl, C_{2-12} alkynyl, C_{3-12} carbocyclyl or C_{3-12} heterocyclyl, wherein the optionally substituted carbocyclyl or heterocyclyl group is optionally fused to one to three unsaturated, partially unsaturated or fully saturated five to seven membered rings containing zero to three heteroatoms;

each substitutable carbon atom in R², including the optional fused ring, is optionally and independently substituted by one or more of C₁₋₁₂ alkyl, C₃₋₁₂ cycloalkyl, C₃₋₁₂ heterocycloalkyl, C₃₋₁₂ aryl, C₃₋₁₂ heteroaryl halogen, C₁₋₁₂ haloalkyl, OR⁹, SR⁹, NO₂, CN, NR⁹R⁹, NR⁹COR⁹, NR⁹COR⁹, NR⁹CO₂R⁹, CO₂R⁹, COR⁹, CONR⁹R⁹, S(O)₂R⁹, SONH₂, S(O)R⁹, SO₂NR⁹R⁹, NR⁹S(O)₂R⁹, wherein each R⁹ may be the same or different and is as defined below and wherein:

the C_{1-12} alkyl group optionally incorporates one or two insertions selected from the group consisting of -O-, -C(O)-, -N(R⁹)-, -S(O)- and -S(O₂)-, wherein each R⁹ may be the same or different and is as defined above;

the C_{1-12} alkyl, C_{3-12} cycloalkyl, C_{3-12} heterocycloalkyl, C_{3-12} aryl, or C_{3-12} heteroaryl groups are optionally substituted by one or more of halogen, C_{1-12} haloalkyl, OR^9 , SR^9 , NO_2 , CN, NR^9R^9 , NR^9COR^9 , NR^9COR^9 , NR^9COR^9 , $NR^9CO_2R^9$, CO_2R^9 , COR^9 , $CONR^9R^9$, $S(O)_2R^9$, $SONH_2$, $S(O)R^9$, $SO_2NR^9R^9$, $NR^9S(O)_2R^9$, wherein each R^9 may be the same or different and is as defined below; and

the C_{3-12} cycloalkyl, C_{3-12} heterocycloalkyl, C_{3-12} aryl, or C_{3-12} heteroaryl groups are optionally substituted by one or more C_{1-12} alkyl groups;

each saturated carbon in R^2 , including the optional fused ring, is further optionally and independently substituted by =O, =S, NNR^9R^9 , =N-OR⁹, =NNHCOR⁹, =NNHCO2R⁹, =NNSO₂R⁹, or =NR⁹, wherein each R⁹ may be the same or different and is as defined below; and

each substitutable nitrogen atom in R² is optionally substituted by R¹⁰, COR⁹, SO₂R⁹ or CO₂R⁹ wherein each R⁹ and R¹⁰ may be the same or different and is as defined below;

wherein two R⁹ in NR⁹₂ may optionally form a partially saturated, unsaturated or fully saturated three to seven membered ring containing one to three heteroatoms, optionally and independently substituted by one or more of C₁₋₆ alkyl, halogen, C₁₋₆ haloalkyl, OR¹¹, SR¹¹, NO₂, CN, NR¹¹R¹¹, NR¹¹COR¹¹, NR¹¹COR¹¹, NR¹¹CO₂R¹¹, CO₂R¹¹, COR¹¹, COR¹¹, COR¹¹₂, S(O)₂R¹¹, SO₂NR¹¹R¹¹, NR¹¹S(O)₂R¹¹,

wherein the C_{1-6} alkyl group optionally incorporates one or two insertions selected from the group consisting of -O-, -N(R¹¹)-, -S(O)- and -S(O₂)-, wherein each R¹¹ may be the same or different and is as defined below;

wherein R¹¹ is hydrogen, C₁₋₆ alkyl, or C₁₋₆ haloalkyl;

wherein R^9 is hydrogen , C_{1-12} alkyl or C_{3-12} aryl, optionally substituted by one or more of C_{1-4} alkyl, halogen, C_{1-4} haloalkyl, OR^{12} , SR^{12} , NO_2 , CN, $NR^{12}R^{12}$, $NR^{12}COR^{12}$, $NR^{12}CONR^{12}R^{12}$, $NR^{12}COR^{12}$, $NR^{12}CO_2R^{12}$, CO_2R^{12} , COR^{12} ,

wherein R^{10} is C_{1-12} alkyl or C_{3-12} aryl, optionally substituted by one or more of C_{1-4} alkyl, halogen, C_{1-4} haloalkyl, OR^{12} , SR^{12} , NO_2 , CN, $NR^{12}R^{12}$, $NR^{12}COR^{12}$, $NR^{12}COR^{12}$, $NR^{12}COR^{12}$, COR^{12} , C

wherein R¹² is hydrogen, C₁₋₄ alkyl, or C₁₋₄ haloalkyl.

4. (Currently Amended) A compound as claimed in any one of claims 1 to 3 claim 1 wherein X is NR³; O, S or (CR²²-R²²)_n wherein R²² is independently one or more of halogen, C₁₋₁₂ aikyi, C₂₋₁₂ aikenyl, C₂₋₁₂ alkynyl, C₁₋₁₂ haloalkyl, C₆₋₁₂ carbocyclyl, C₅₋₁₂ heterocyclyl, (CH₂)_nOR⁵, (CH₂)_nNR⁵₂, OR⁵, SR⁵, NO₂, CN, NR⁵₂, NR⁵COR⁵, NR⁵CONR⁵₂, NR⁵COR⁵, NR⁵CO₂R⁵, CO₂R⁵, CO₂R⁵, COR⁵, CONR⁵₂, S(O)₂R⁵, SONR⁵₂, S(O)₂R⁵, SO₂NR⁵₂, or NR⁵S(O)₂R⁵ wherein each R⁵ may be the same or different and is as defined above; and

wherein n is 1 to 6, preferably n is 1, 2, 3 or 4;

and wherein R^3 is a lone electron pair, hydrogen, C_{1-12} alkyl, C_{2-12} alkenyl, C_{2-12} alkynyl, C_{3-12} carbocyclyl or C_{3-12} heterocyclyl, each of which is optionally substituted, wherein:

the optionally substituted carbocyclyl or heterocyclyl group is optionally fused to one to three unsaturated, partially unsaturated or fully saturated five to seven membered rings containing zero to three heteroatoms,

each substitutable carbon atom in R^3 , including the optional fused ring, is optionally and independently substituted by one or more of C_{1-12} alkyl, C_{3-12} cycloalkyl, C_{3-12} heterocycloalkyl, C_{3-12} aryl, C_{3-12} heteroaryl halogen, C_{1-12} haloalkyl, OR^{13} , SR^{13} , NO_2 , CN, $NR^{13}R^{13}$, $NR^{13}COR^{13}$, $NR^{13}COR^{13}$, $NR^{13}COR^{13}$, $NR^{13}COR^{13}$, $NR^{13}COR^{13}$, $NR^{13}CO_2R^{13}$, CO_2R^{13} , COR^{13} ,

the C_{1-12} alkyl group optionally incorporates one or two insertions selected from the group consisting of -O-, -C(O)-, -N(R¹³)-, -S(O)- and -S(O₂)-, wherein each R¹³ may be the same or different and is as defined above:

the C_{1-12} alkyl, C_{3-12} cycloalkyl, C_{3-12} heterocycloalkyl, C_{3-12} aryl, or C_{3-12} heteroaryl groups are optionally substituted by one or more of halogen, C_{1-12} haloalkyl, OR^{13} , SR^{13} , NO_2 , CN, $NR^{13}R^{13}$, $NR^{13}COR^{13}$, $NR^{13}COR^{13}$, $NR^{13}COR^{13}$, $NR^{13}CO_2R^{13}$, CO_2R^{13} , CO_2R^{13} , CO_3R^{13} , $CO_3R^$

CONR¹³R¹³, S(O)₂R¹³, SONH₂, S(O)R¹³, SO₂NR¹³R¹³, NR¹³S(O)₂R¹³, wherein each R¹³ may be the same or different and is as defined below; and

the C_{3-12} cycloalkyl, C_{3-12} heterocycloalkyl, C_{3-12} aryl, or C_{3-12} heteroaryl groups are optionally substituted by one or more C_{1-12} alkyl groups;

each saturated carbon in R^2 , including the optional fused ring, is further optionally and independently substituted by =0, =S, $NNR^{13}R^{13}$, =N-OR¹³, =NNHCOR¹³, =NNHCO₂R¹³, =NNHCO₂R¹³, or =NR¹³, wherein each R^{13} may be the same or different and is as defined below; and

each substitutable nitrogen atom in R^3 is optionally substituted by R^{14} , COR^{13} , SO_2R^{13} or CO_2R^{13} wherein each R^{13} and R^{14} may be the same or different and is as defined below;

wherein two R¹³ in NR¹³₂ may optionally form a partially saturated, unsaturated or fully saturated three to seven membered ring containing one to three heteroatoms, optionally and independently substituted by one or more of C₁₋₆ alkyl, halogen, C₁₋₆ haloalkyl, OR¹⁵, SR¹⁵, NO₂, CN, NR¹⁵R¹⁵, NR¹⁵COR¹⁵, NR¹⁵COR¹⁵, NR¹⁵COR¹⁵, NR¹⁵CO₂R¹⁵, CO₂R¹⁵, COR¹⁵, COR¹⁵, COR¹⁵, S(O)₂R¹⁵, SONR¹⁵₂, S(O)_R¹⁵, SO₂NR¹⁵R¹⁵, NR¹⁵S(O)₂R¹⁵,

wherein the C_{1-6} alkyl group optionally incorporates one or two insertions selected from the group consisting of -O-, -N(R^{15})-, -S(O)- and -S(O₂)-, wherein each R^{15} may be the same or different and is as defined below;

wherein R¹⁵ is hydrogen, C₁₋₆ alkyl, or C₁₋₆ haloalkyl;

wherein R^{13} is hydrogen , C_{1-12} alkyl or C_{3-12} aryl, optionally substituted by one or more of C_{1-4} alkyl, halogen, C_{1-4} haloalkyl, OR^{16} , SR^{16} , NO_2 , CN, $NR^{16}R^{16}$, $NR^{16}COR^{16}$, $NR^{16}CONR^{16}R^{16}$, $NR^{16}COR^{16}$, $NR^{16}CO_2R^{16}$, CO_2R^{16} , CO_2R^{16} , COR^{16} , $CONR^{16}_2$, $S(O)_2R^{16}$, $SONH_2$, $S(O)R^{16}$, SO_2 $NR^{16}R^{16}$, $NR^{16}S(O)_2R^{16}$, wherein the C_{1-12} alkyl group optionally incorporates one or two insertions selected from the group consisting of -O-, -N(R^{16})-, -S(O)- and -S(O₂)-, wherein each R^{16} may be the same or different and is as defined below;

wherein R^{14} is C_{1-12} alkyl or C_{3-12} aryl, optionally substituted by one or more of C_{1-4} alkyl, halogen, C_{1-4} haloalkyl, OR^{16} , SR^{16} , NO_2 , CN, $NR^{16}R^{16}$, $NR^{16}COR^{16}$, $NR^{16}CONR^{16}R^{16}$, $NR^{16}COR^{16}$, $NR^{16}CO_2R^{16}$, CO_2R^{16} , CO_2R^{16} , COR^{16} , $CONR^{16}_2$, $S(O)_2R^{16}$, $SONH_2$, $S(O)R^{16}$, $SO_2NR^{16}R^{16}$, $NR^{16}S(O)_2R^{16}$, wherein the C_{1-12} alkyl group optionally incorporates one or two insertions selected from the group consisting of -O-, -N(R^{16})-, -S(O)- and -S(O₂)-, wherein each R^{16} may be the same or different and is as defined below;

wherein R¹⁶ is hydrogen, C₁₋₄ alkyl, or C₁₋₄ haloalkyl;

wherein when X is NR², R² and R³ can form a 3 to 12 membered heterocyclyl ring, more preferably a 5, 6, 7, 8, 9, 10, 11 or 12 membered ring, wherein said ring can be partially saturated, unsaturated or fully saturated containing one to three heteroatoms; wherein the heterocyclylic group formed by R² and R³ can be optionally fused to one to three unsaturated, partially saturated or fully saturated 5 to 7 membered rings containing zero to three heteroatoms, any of said rings being optionally and independently substituted with one or more of C₁₋₆ alkyl, halogen, C₁₋₆ haloalkyl, OR²², SR²², NO₂, CN, NR²²R²², NR²²COR²², NR²²CONR²²R²², NR²²COR²², CO₂R²², CO₂R²², COR²², CONR²²₂, S(O)₂R²², SONR²²₂, S(O)₂R²², SONR²²₂, S(O)₂R²², SO₂NR²²R²², NR²²S(O)₂R²², wherein the C₁₋₆ alkyl group optionally incorporates one or two insertions from -O-, -N(R²²)-, -S(O)- and -S(O₂)- and wherein each R²² may be the same or different.

5. (Currently Amended) A compound as claimed in any one of claims 1 to 4 claim 1 wherein R⁴ is a six-membered carbocyclyl group or a five or six-membered heterocyclyl group containing from 1 to 4 heteroatoms independently selected from N, S or O, wherein the optionally substituted six-membered carbocyclyl or five or six-membered heterocyclyl group is optionally fused to a partially saturated, unsaturated or fully saturated five to seven membered ring containing zero to three heteroatoms, and each substitutable carbon or hetero-atom in R⁴ including the optional fused ring, is optionally and independently substituted by one or more of halogen, C₁₋₁₂ alkyl, C₂₋₁₂ alkenyl, C₂₋₁₂ alkynyl, C₁₋₁₂ haloalkyl, C₃₋₁₂ carbocyclyl, C₃₋₁₂ heterocyclyl, (CH₂)_nOR¹⁷, (CH₂)_nNR¹⁷₂, OR¹⁷, SR¹⁷, NO₂, CN, NR¹⁷₂, NR¹⁷COR¹⁷, NR¹⁷CONR¹⁷₂, NR¹⁷COR¹⁷, CO₂R¹⁷, CO₂R¹⁷, COR¹⁷, CONR¹⁷₂, S(O)₂R¹⁷, SONR¹⁷₂, S(O)₂R¹⁷, SONR¹⁷₂, S(O)₂R¹⁷, SONR¹⁷₂, S(O)₂R¹⁷, SONR¹⁷₂, S(O)₂R¹⁷, SONR¹⁷₂, S(O)₂R¹⁷, wherein the C₁₋₁₂ alkyl group optionally contains one or

more insertions selected from -O-, -N(R¹²)- -S-, -S(O)- and -S(O₂)-; and each saturated carbon in the optional fused ring is further optionally and independently substituted by =O, =S, NNR¹⁸₂, =N-OR¹⁸, =NNR¹⁸CO2¹⁸, =NNR¹⁸CO2¹⁸, =NNSO2¹⁸, or =NR¹⁸; and each substitutable nitrogen atom in R⁴ is optionally substituted by R¹⁹, COR¹⁹, SO2¹⁹ or CO2¹⁹; wherein n is 1 to 6, preferably n is 1, 2 or 3; preferably, wherein each substitutable carbon or hetero-atom in R⁴ is optionally and independently substituted by one or more of C₁₋₆ alkyl, OR²⁰, SR²⁰, NO₂, CN, NR²⁰₂, NR²⁰COR²⁰, NR²⁰COR²⁰, NHCO2²⁰, CO2²⁰, CO2²⁰, COR²⁰, CONR²⁰₂, S(O)2²⁰, SONR²⁰₂, S(O)R²⁰, SO2NR²⁰₂, or NR²⁰S(O)2²⁰;

wherein R²⁰ is hydrogen, C₁₋₆ alkyl, or C₁₋₆ haloalkyl;

wherein R^{17} is hydrogen , C_{1-12} alkyl, C_{3-12} carbocyclyl or C_{3-12} heterocyclyl, optionally substituted by one or more of C_{1-6} alkyl, C_{3-12} carbocyclyl, C_{3-12} heterocyclyl, halogen, C_{1-6} haloalkyl, OR^{21} , SR^{21} , NO_2 , CN, $NR^{21}R^{21}$, $NR^{21}COR^{21}$, $NR^{21}CONR^{21}R^{21}$, $NR^{21}COR^{21}$, $NR^{21}COR^{21}$, $NR^{21}CO_2R^{21}$, CO_2R^{21} , CO_2R^{21} , COR^{21} , CO

wherein two R^{17} in NR^{17}_2 may optionally form a partially saturated, unsaturated or fully saturated three to seven membered ring containing one to three heteroatoms, optionally and independently substituted by one or more of C_{1-6} alkyl, halogen, C_{1-6} haloalkyl, OR^{21} , SR^{21} , NO_2 , CN, $NR^{21}R^{21}$, $NR^{21}COR^{21}$, $NR^{21}COR^{21}$, $NR^{21}COR^{21}$, $NR^{21}CO_2R^{21}$, CO_2R^{21} , $CO_2R^{$

wherein R^{18} is hydrogen, C_{1-12} alkyl, C_{3-12} carbocyclyl or C_{3-12} heterocyclyl, optionally substituted by one or more of C_{1-6} alkyl, halogen, C_{1-6} haloalkyl, OR^{21} , SR^{21} , NO_2 , CN, $NR^{21}R^{21}$, $NR^{21}COR^{21}$, wherein the C_{1-12} alkyl group optionally incorporates one

or two insertions selected from the group consisting of -O-, -N(R^{21})-, -S(O)- and -S(O₂)-, wherein each R^{21} may be the same or different and is as defined below;

wherein R^{19} is hydrogen, C_{6-12} aryl, C_{1-6} alkyl or C_{1-6} haloalkyl;

wherein R²¹ is hydrogen, C₁₋₆ alkyl, or C₁₋₆ haloalkyl.

- 6. (Currently Amended) A compound as claimed in any one of claims 1 to 5 claim 1 wherein R¹ is an optionally substituted five or six membered carbocyclyl or heterocyclyl group selected from optionally substituted phenyl, acridine, benzimidazole, benzofuran, benzothiophene, benzoxazole, benzothiazole, cyclohexyl furan, imidazole, indole, isoindole, isoquinoline, isoxazole, isothiazole, morpholine, napthaline, oxazole, phenazine, phenothiazine, phenoxazine, piperazine, piperidine, pyrazole, pyridazine, pyridine, pyrrole, quinoline, quinolizine, tetrahydrofuran, tetrazine, tetrazole, thiophene, thiazole, thiomorpholine, thianaphthalene, thiopyran, triazine, triazole or trithiane.
- 7. (Currently Amended) A compound as claimed in any one of claims 1 to 6 claim 1 wherein R¹ is a group of formula (II), wherein X is a group NR³, Y is absent and one or more of R² and R³ is hydrogen, alkyl or cycloalkyl.
- 8. (Original) A compound as claimed in claim 7 wherein the group of formula (II) is an alkylamino or cycloalkylamino group preferably selected from optionally substituted methylamino, ethylamino, propylamino, isopropylamino, butylamino, cyclobutylamino, pentylamino, cyclopentylamino, hexylamino, cyclohexylamino, heptylamino, cycloheptylamino, octylamino and cyclooctylamino.
- 9. (Currently Amended) A compound as claimed in any one of claims 1 to 8 claim 1 wherein R¹ is substituted with one or more of OR²⁴, halogen, C₁₋₆ alkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl, C₁₋₆haloalkyl, C₁₋₆alkylaryl, C₁₋₆alkylheterocyclyl, (CH₂)_nOR²⁴, (CH₂)_nNR²⁴₂, SR²⁴, NO₂, CN, NR²⁴₂, CO₂R²⁴, NR²⁴C(O)R²⁴, NR²⁴S(O)₂R²⁴, COR²⁴, CONR²⁴₂, S(O)₂R²⁴, S(O)₂R²⁴ or SO₂NR²⁴₂;

wherein R^{24} is hydrogen, C_{1-4} alkyl or C_{6-12} aryl preferably phenyl, or C_{5-12} heterocyclyl preferably pyridine, and n is 1, 2, 3, 4, 5 or 6;

wherein two R²⁴ in NR²⁴₂ may optionally form a partially saturated, unsaturated or fully saturated three to seven membered ring containing one to three heteroatoms, said ring is preferably independently substituted with one or more of halogen, C₁₋₁₂ alkyl, C₂₋₁₂ alkenyl, C₂₋₁₂ alkynyl, C₁₋₁₂ haloalkyl, C₃₋₁₂ carbocyclyl, C₃₋₁₂ heterocyclyl, OR²⁵, SR²⁵, NO₂, CN, NR²⁵₂, NR²⁵COR²⁵, NR²⁵COR²⁵, NR²⁵COR²⁵, NR²⁵CO₂R²⁵, CO₂R²⁵, COR²⁵, CONR²⁵₂, S(O)₂R²⁵, SONR²⁵₂, S(O)₂R²⁵₂, or NR²⁵S(O)₂R²⁵; and each saturated carbon in the optional ring is further optionally and independently substituted by =O, =S, NNR²⁶₂, =N-OR²⁶, =NNR²⁶COR²⁶, =NNR²⁶COR²⁶, or =NR²⁶; and each substitutable nitrogen atom is optionally substituted by R²⁷, COR²⁷, SO₂R²⁷ or CO₂R²⁷;

wherein R^{25} is hydrogen , $C_{1\text{-}12}$ alkyl, $C_{6\text{-}12}$ carbocyclyl or $C_{5\text{-}12}$ heterocyclyl, optionally substituted by one or more of $C_{1\text{-}6}$ alkyl, halogen, $C_{1\text{-}6}$ haloalkyl, OR^{28} , SR^{28} , NO_2 , CN, $NR^{28}R^{28}$, $NR^{28}COR^{28}$, $NR^{28}COR^{28}$, $NR^{28}COR^{28}$, $NR^{28}CO_2R^{28}$, CO_2R^{28} , CO_2R^{28} , COR^{28} , CO

wherein R^{26} is hydrogen, C_{1-12} alkyl, C_{6-12} carbocyclyl or C_{5-12} heterocyclyl, optionally substituted by one or more of C_{1-6} alkyl, halogen, C_{1-6} haloalkyl, OR^{28} , SR^{28} , NO_2 , CN, $NR^{28}R^{28}$, $NR^{28}COR^{28}$, $NR^{28}COR^{28}$, $NR^{28}COR^{28}$, $NR^{28}COR^{28}$, $NR^{28}CO_2R^{28}$, CO_2R^{28} , CO_2R^{28} , COR^{28} ,

wherein R^{27} is hydrogen, C_{1-6} alkyl, C_{1-6} haloalkyl or C_{6-12} aryl;

wherein R²⁸ is hydrogen, C₁₋₆ alkyl, or C₁₋₆ haloalkyl.

10. (Currently Amended) A compound as claimed in any one of claims 1 to 9 claim 1 wherein R⁴ is selected from phenyl, cyclohexyl, acridine, benzimidazole, benzofuran,

benzothiophene, benzoxazole, benzothiazole, indole, isoindole, indolizine, indazole, isoindole, isoquinoline, morpholine, napthalene, phenazine, phenothiazine, phenoxazine, piperazine, piperidine, pyridazine, pyridine, pyridinyl, pyrimidinyl, pyrazinyl, pyridazinyl, quinoline, quinolizine, tetrazine, thiomorpholine, thianaphthalene, thiopyran, triazine, trithiane, furan, imidazole, isoxazole, isothiazole, oxazole, oxadiazole, oxathiazole, pyrazole, pyrrole, tetrazole, thiophene, thiadiazole, thiatriazole, thiazole or triazole, wherein each substitutable carbon or hetero-atom in R⁴ is optionally and independently substituted by one or more of C₁₋₆ alkyl, OR²⁰, SR²⁰, NO₂, CN, NR²⁰₂, NR²⁰COR²⁰, NR²⁰CONR²⁰₂, NR²⁰COR²⁰, NHCO₂R²⁰, CO₂R²⁰, COR²⁰, CONR²⁰₂, S(O)₂R²⁰, SONR²⁰₂, S(O)₂R²⁰, SONR²⁰₂, S(O)₂R²⁰, OR²⁰₂, OR²⁰₂,

wherein R²⁰ is hydrogen, C₁₋₆ alkyl, or C₁₋₆ haloalkyl.

11. (Currently Amended) A compound as claimed in any one of claims 1 to 10 claim 1 wherein R^4 is a six-membered carbocyclyl or heterocyclyl group optionally substituted with one or more of OR^{29} , NR^{29}_2 , SR^{29} , $(CH_2)_nOR^{29}$, $(CH_2)_nNR^{29}_2$, halogen, C_{1-6} alkyl, C_{2-6} alkenyl, C_{2-6} alkynyl, haloalkyl, NO_2 , CN, $NR^{29}C(O)R^{29}$, $NR^{29}S(O)_2R^{29}$, CO_2R^{29} , COR^{29} , $CONR^{29}_2$, $S(O)_2R^{29}$, $S(O)_2R^{29}$, $S(O)_2R^{29}_2$;

wherein R^{29} is hydrogen, C_{1-4} alkyl, C_{5-12} heterocyclyl or C_{6-12} aryl preferably phenyl, and n is 1, 2, 3, 4, 5 or 6;

wherein two R^{29} in NR^{29}_2 may optionally form a partially saturated, unsaturated or fully saturated five to seven membered ring containing one to three heteroatoms, optionally and independently substituted with one or more of halogen, C_{1-12} alkyl, C_{2-12} alkenyl, C_{2-12} alkynyl, C_{1-12} haloalkyl, C_{6-12} carbocyclyl, C_{5-12} heterocyclyl, OR^{30} , SR^{30} , NO_2 , CN, NR^{30}_2 , $NR^{30}COR^{30}$, $NR^{30}COR^{30}$, $NR^{30}COR^{30}$, $NR^{30}COR^{30}$, $NR^{30}CO_2R^{30}$, CO_2R^{30} , COR^{30} , $CONR^{30}_2$, $S(O)_2R^{30}$, $SO_2NR^{30}_2$, or $NR^{30}S(O)_2R^{30}$; and each saturated carbon in the optional ring is further optionally and independently substituted by =O, =S, NNR^{31}_2 , $=N-OR^{31}$, $=NNR^{31}COR^{31}$, $=NNR^{31}CO_2R^{31}$, $=NNSO_2R^{31}$, or $=NR^{31}$; and each substitutable nitrogen atom is optionally substituted by R^{32} , COR^{32} , SO_2R^{32} or CO_2R^{32} ;

wherein R^{30} is hydrogen , C_{1-12} alkyl, C_{6-12} carbocyclyl or C_{5-12} heterocyclyl, optionally substituted by one or more of C_{1-6} alkyl, halogen, C_{1-6} haloalkyl, OR^{33} , SR^{33} , NO_2 , CN, $NR^{33}R^{33}$,

 $NR^{33}COR^{33}$, $NR^{33}CONR^{33}R^{33}$, $NR^{33}COR^{33}$, $NR^{33}CO_2R^{33}$, CO_2R^{33} , COR^{33} , $CONR^{33}_2$, $S(O)_2R^{33}$, $SONR^{33}_2$, $S(O)_2R^{33}$, $SO_2NR^{33}R^{33}$, wherein the C_{1-12} alkyl group optionally incorporates one or two insertions selected from the group consisting of -O-, -N(R^{33})-, -S(O)- and -S(O₂)-, wherein each R^{33} may be the same or different and is as defined below;

wherein R^{31} is hydrogen, C_{1-12} alkyl, C_{6-12} carbocyclyl or C_{5-12} heterocyclyl, optionally substituted by one or more of C_{1-6} alkyl, halogen, C_{1-6} haloalkyl, OR^{33} , SR^{33} , NO_2 , CN, $NR^{33}R^{33}$, $NR^{33}COR^{33}$, $NR^{33}COR^{33}$, $NR^{33}COR^{33}$, $NR^{33}COR^{33}$, $NR^{33}COR^{33}$, $NR^{33}CO_2R^{33}$, CO_2R^{33} , COR^{33} , COR^{3

wherein R^{32} is hydrogen, C_{6-12} aryl, C_{1-6} alkyl or C_{1-6} haloalkyl;

wherein R³³ is hydrogen, C₁₋₆ alkyl, or C₁₋₆ haloalkyl.

12. (Currently Amended) A compound as claimed in any one of claims 1 to 11 claim 1 wherein R⁴ is a five-membered heterocyclyl,

wherein A, X^2 , Y^2 or Z are independently selected from N, O, C, S and M is C or N, wherein one, two, three or four of A, X^2 , Y^2 , Z and M is other than C;

 R^{34} , R^{35} , R^{36} or R^{37} are independently selected from a lone electron pair, hydrogen, halogen, C_{1-12} alkyl, C_{1-12} haloalkyl, OR^{38} , SR^{38} , NO_2 , CN, NR^{38}_2 , $NR^{38}COR^{38}$, $NR^{38}CONR^{38}_2$, $NR^{38}COR^{38}$, $NR^{38}COR^{38}_2$

wherein n is 1 to 6, preferably n is 1, 2 or 3;

or wherein any two of R³⁴, R³⁵, R³⁶ or R³⁷ may optionally form a partially saturated, unsaturated or fully saturated five to seven membered ring containing zero to three heteroatoms, each saturated carbon in the optional fused ring is further optionally and independently substituted with one or more of halogen, C₁₋₁₂ alkyl, C₂₋₁₂ alkenyl, C₂₋₁₂ alkynyl, C₁₋₁₂ haloalkyl, C₆₋₁₂ carbocyclyl, C₅₋₁₂ heterocyclyl, OR³⁸, SR³⁸, NO₂, CN, NR³⁸₂, NR³⁸CONR³⁸₂, NR³⁸COR³⁸, NR³⁸CO₂R³⁸, (CH₂)_nOR³⁸, (CH₂)_nNR³⁸₂, CO₂R³⁸, COR³⁸, CONR³⁸₂, S(O)₂R³⁸, SONR³⁸₂, S(O)₂R³⁸, SONR³⁸₂, S(O)₂R³⁸, and each saturated carbon in the optional fused ring is further optionally and independently substituted by =O, =S, NNR³⁹₂, =N-OR³⁹, =NNR³⁹COR³⁹, =NNR³⁹CO₂R³⁹, or =NR³⁹; and each substitutable nitrogen atom in R⁴ is optionally substituted by R⁴⁰, COR⁴⁰, SO₂R⁴⁰ or CO₂R⁴⁰;

wherein n is 1 to 6, preferably n is 1, 2 or 3;

wherein R^{38} is hydrogen, C_{1-12} alkyl, C_{6-12} carbocyclyl or C_{5-12} heterocyclyl, optionally substituted by one or more of C_{1-6} alkyl, halogen, C_{1-6} haloalkyl, OR^{41} , SR^{41} , NO_2 , CN, $NR^{41}R^{41}$, $NR^{41}CONR^{41}R^{41}$, $NR^{41}COR^{41}$, $NR^{41}CO_2R^{41}$, CO_2R^{41} , COR^{41} , $CONR^{41}_2$, CO

wherein R^{39} is hydrogen, C_{1-12} alkyl, carbocyclyl or heterocyclyl, optionally substituted by one or more of C_{1-6} alkyl, halogen, C_{1-6} haloalkyl, OR^{41} , SR^{41} , NO_2 , CN, $NR^{41}R^{41}$, $NR^{41}COR^{41}$, $NR^{41}COR^{41}$, $NR^{41}COR^{41}$, $NR^{41}COR^{41}$, $NR^{41}COR^{41}$, $NR^{41}CO_2R^{41}$, CO_2R^{41} , COR^{41} , $CONR^{41}_2$, $S(O)_2R^{41}$, $S(O)_2R^{41}$, $SO_2NR^{41}R^{41}$, $NR^{41}S(O)_2R^{41}$, wherein the C_{1-12} alkyl group optionally incorporates one or two insertions selected from the group consisting of -O-, $-N(R^{41})$ -, -S(O)- and $-S(O_2)$ -, wherein each R^{41} may be the same or different and is as defined below;

wherein R^{40} is hydrogen, C_{1-6} alkyl, C_{1-6} haloalkyl or C_{6-12} aryl.

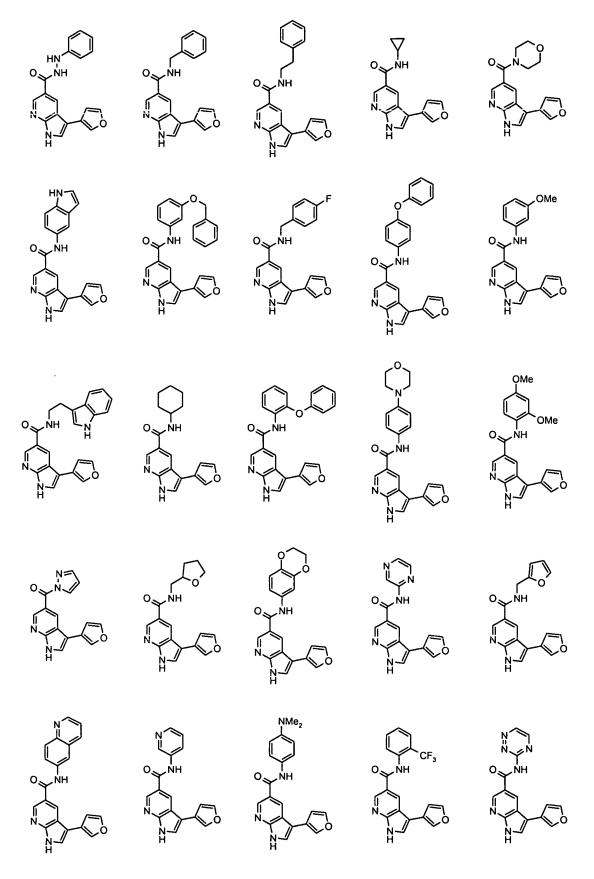
wherein R⁴¹ is hydrogen, C₁₋₆ alkyl, or C₁₋₆ haloalkyl.

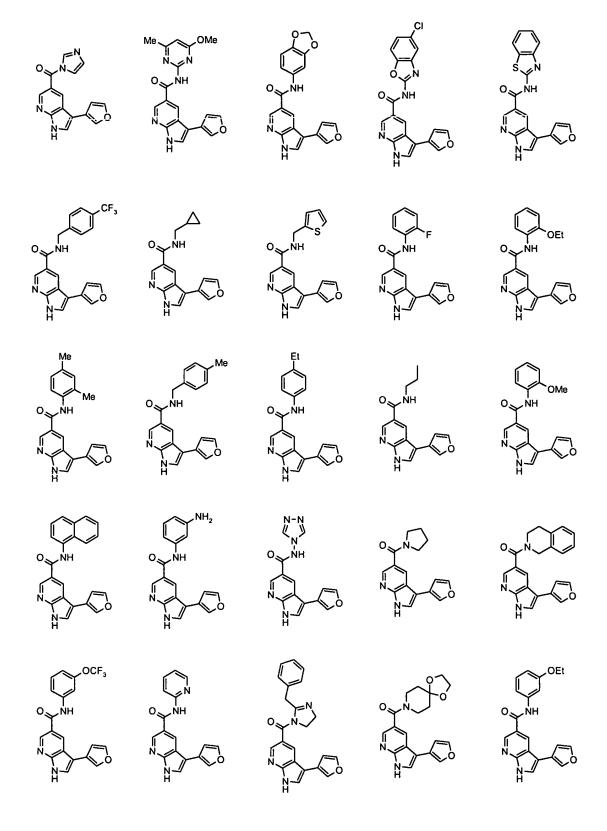
13. (Original) A compound as claimed in claim 12 wherein R⁴ is furan, imidazole, isoxazole, isothiazole, oxazole, oxadiazole, oxatriazole, pyrazole, pyrrole, tetrazole, thiophene, thiadiazole,

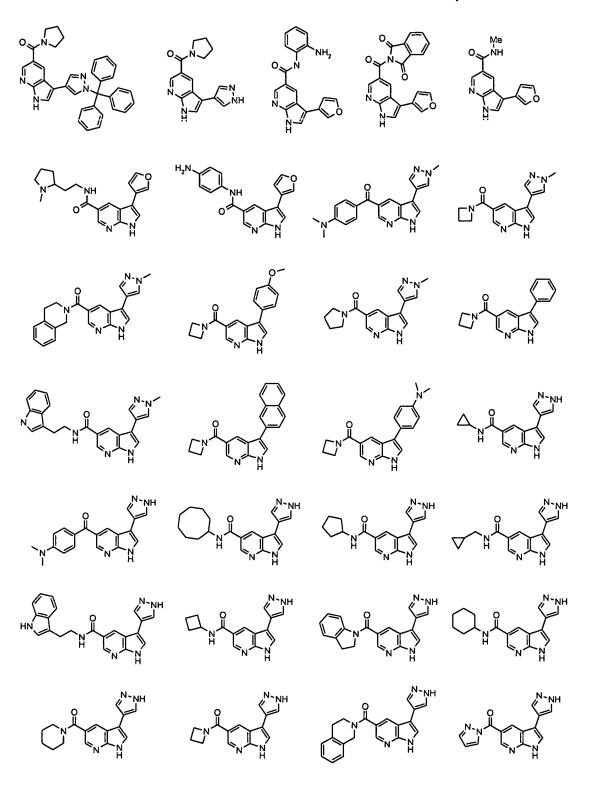
thiatriazole, thiazole or triazole; and R^{34} , R^{35} , R^{36} or R^{37} are independently selected from a lone electron pair, hydrogen, halogen, C_{1-6} alkyl, C_{1-6} haloalkyl, OR^{42} , SR^{42} , CN, NR^{42}_{2} , $NR^{42}COR^{42}$, CO_2R^{42} , COR^{42} ,

wherein R⁴² is hydrogen, C₁₋₄ alkyl, preferably methyl or ethyl or carbocyclyl, preferably phenyl.

14. (Currently Amended) A compound as claimed in any one of claims 1 to 13 claim 1 selected from the group consisting of:







15. (Currently Amended) A process for the manufacture of a compound of formula (I) wherein R¹ is a group of formula (II) as defined in the any one of claims 1 to 14 claim 1 of the invention comprising the condensation of an intermediate (III) with an intermediate (IV).

wherein R^2 and R^4 are as defined in any one of claims 1 to 14; and wherein each of L^1 and L^2 is independently a leaving group wherein L^1 and L^2 together form a condensation product.

- 16. (Original) A process as claimed in claim 15 wherein L^1 is OH, OR^{50} , OM, Cl, Br or I wherein R^{50} is C_{1-6} alkyl, preferably methyl or ethyl and M is Na, Li, K, Ca, Mg or Ba, and L^2 is hydrogen or M.
- 17. (Original) A compound of formula (III)

wherein R⁴ is as defined any one of claims 1 to 14

$$R^{50}$$
 is C_{1-6} alkyl, and

M is Na, Li, K, Ca, Mg, or Ba.

18. (Original) A process for the manufacture of a compound of formula (V) comprising removal of group R⁵¹ from an intermediate (VI)

wherein L^3 is R^1 or L^1 ;

R¹ and R⁴ are as defined in any one of claims 1 to 14;

L¹ is as defined in claim 17;

and R⁵¹ is an amino protecting group selected from R⁵²SO₂, R⁵²C(O), R⁵²₃Si, R⁵²OCH₂, (R⁵²)₂NSO₂, (R⁵²)₂NC(O), R⁵²OC(O), R⁵²(R⁵²O)CH, R⁵²CH₂CH₂, R⁵²CH₂, PhC(O)CH₂, CH₂=CH, ClCH₂CH₂, Ph₃C, Ph₂(4-pyridyl)C, Me₂N, HO-CH₂, R⁵²OCH₂, (R⁵²)₃SiOCH₂, (R⁵²O)₂CH, t-BuOC(O)CH₂, Me₂NCH₂, and tetrahydropyranylamine,

wherein R^{52} is C_{1-6} alkyl or C_{6-12} aryl.

19. (Original) A compound of formula (VI)

wherein R⁴ is as defined in any one of claims 1 to 14, and

wherein L^3 and R^{51} are as defined in claim 18.

20. (Currently Amended) A process for the manufacture of a compound of formula (VI) comprising a a) a reaction of a compound of formula (VII) with stannane R^4 -Sn(R^{53})₃ in the presence of a palladium catalyst or b) reaction of a compound of formula (VII) with boronic acid or ester R^4 -B(OR⁵⁴)₂ in a presence of a suitable palladium catalyst or c) reaction of a compound of formula (VII) with silane R^4 -Si(R^{55})₃ in the presence of a palladium catalyst;

wherein R⁴ is as defined in any one of claims 1 to 14,

L³ is as defined in claim 18;

R⁵¹ is an amino protecting group as defined in claim 18;

X³ is F, Cl, Br I or CF₃SO₃,

and R^{53} is independently C_{1-6} alkyl;

 R^{54} is independently hydrogen or C_{1-6} alkyl or wherein two R^{54} groups together optionally form a five, six or seven membered ring with the boron and oxygen atoms, wherein the ring is optionally substituted with one or more C_{1-6} alkyl group.

and R^{55} is independently $C_{1\text{-}6}$ alkyl, F, or OH.

- 21. (Original) A process as claimed in claim 20 wherein the catalyst is one or more selected from $(PPh_3)_2PdCl_2$, $(PPh_3)_4Pd$, $Pd(OAc)_2$, $[PdCl(\eta^3-C_3H_5]_2$, $Pd_2(dba)_3$, $Pd(dba)_2$ (dba = dibenzylidenacetone), and $Pd/P(t-Bu)_3$.
- 22. (Original) A compound of formula (VII)

wherein L³ is as defined in claim 18;

wherein R⁵¹ is an amino protecting group as defined in claim 18;

wherein X^3 is as defined in claim 20.

23. (Original) A process for the manufacture of a compound of formula (VII) comprising protection of the pyrrole nitrogen with a group R⁵¹,

wherein L³ is as defined in claim 18;

wherein R⁵¹ is an amino protecting group defined in claim 18;

wherein X^3 is as defined in claim 20.

24. (Original) A compound of formula (VIII)

wherein L³ is as defined in claim 18;

and X³ is as defined in claim 20.

25. (Original) A process for the production of a compound of formula (VIII) by the introduction of an X³ group into a compound of formula (IX)

wherein L³ is as defined in claim 18 and X³ is as defined in claim 20.

26. (Currently Amended) A compound of formula (IX)

wherein L^3 is a group L^1 as defined in claim 17 or a group R^1 , wherein R^1 is a group of formula (II)

wherein X is NR³, O, S or $(CR^{22}R^{22})_n$, Y is absent or is NR²³, O, or $(CR^{23}R^{23})_n$, R² is optionally substituted C_{1-12} alkyl, C_{2-12} alkenyl, C_{2-12} alkynyl, C_{3-12} carbocyclyl or C_{3-12} heterocyclyl as elaimed defined in any one of claims 1 to 14.

27. (Original) A process for the production of a compound of formula (VII) by the introduction of a X³ group to a compound of formula (X)

wherein L³ and R⁵¹ are as defined in claim 18 and X³ is as defined in claim 20.

28. (Currently Amended) A compound of formula (X)

wherein L^3 is a group L^1 as defined in claim 17 or a group R^1 , wherein R^1 is a group of formula (II)

wherein X is NR³, O, S or $(CR^{22}R^{22})_n$, Y is absent or is NR²³, O, or $(CR^{23}R^{23})_n$, R² is optionally substituted C_{1-12} alkyl, C_{2-12} alkenyl, C_{2-12} alkynyl, C_{3-12} carbocyclyl or C_{3-12} heterocyclyl as elaimed defined in any one of claims 1 to 14;

and R⁵¹ is an amino protecting group as defined in claim 18.

29. (Original) A process for the preparation of compound of formula (IX) by the acidcatalysed hydrolysis of nitrile (XI) in the presence of an alcohol,

$$\begin{array}{cccc}
CN & & & & & & & & & \\
N & & & & & & & & & & \\
N & & & & & & & & & \\
N & & & & & & & & & \\
N & & & & & & & & \\
N & & & & & & & & \\
N & & & & & & & \\
N & & & & & & & \\
N & & & & & & & \\
N & & & & & & & \\
N & & & & \\
N & & & & \\$$

wherein L^3 is OR^{50} ;

and R⁵⁰ is as defined in claim 16.

30. (Original) A compound of formula (XI)

$$\widetilde{\mathbf{z}}$$
 $\widetilde{\mathbf{z}}$ $\widetilde{\mathbf{z}}$

31. (Original) A process for the production of 1H-Pyrrolo[2,3-b]pyridine-5-carbonitrile (XI) comprising reaction of 5-bromo-1*H*-pyrrolo[2,3-*b*]pyridine with Zn(CN)₂ in the presence of a palladium catalyst.

$$\begin{array}{c|c}
Br & CN \\
\hline
N & Pd catalyst & N \\
\end{array}$$
(XI)

32. (Currently Amended) A compound as claimed in any one of claims 17, 19, 22, 24, 26, 28 or 30 selected from one or more of the group consisting of:

- 33. (Canceled)
- 34. (Currently Amended) A pharmeceutical composition comprising a compound as claimed in any one of claims 1 to 14 claim 1 in combination with a pharmaceutically acceptable carrier, diluent or excipient.
- 35. (Original) A composition as claimed in claim 34, additionally comprising one or more of an anti-inflammatory agent, an AMPA receptor antagonist, a chemotherapeutic agent and/or an antiproliferative agent.
- 36. (Canceled)
- 37. (Canceled)
- 38. (Canceled)
- 39. (Canceled)
- 40. (Canceled)
- 41. (Canceled)

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42. (Canceled)

43. (Canceled)

44. (Canceled)

45. (Canceled)

46. (Currently Amended) A method of treating or preventing a -mediated disorder in an

individual, which method comprises administering to said individual a compound as claimed in

any of claims 1-14 claim 1 or a composition as claimed in any of claims 34 or 35.

47. (Canceled)

48. (Currently Amended) A method as claimed in claim 46 or 47, wherein the disorder is a

neurodegenerative disorder, inflammatory disease, a disorder linked to apoptosis, particularly

neuronal apoptosis, autoimmune disease, destructive bone disorder, proliferative disorder,

cancer, infectious disease, allergy, ischemia reperfusion injury, heart attack, angiogenic disorder,

organ hypoxia, vascular hyperplasia, cardiac hypertrophy, thrombin induced platelet aggregation

and/or any condition associated with prostaglandin endoperoxidase synthase-2.

49. (Original) A method as claimed in claim 48, wherein the neurodegenerative disorder

results from apoptosis and/or inflammation.

50. (Currently Amended) A method as claimed in claim 48 or 49, wherein the

neurodegenerative disorder is: dementia; Alzheimer's disease; Parkinson's disease; Amyotrophic

Lateral Sclerosis; Huntington's disease; senile chorea; Sydenham's chorea; hypoglycemia; head

and spinal cord trauma including traumatic head injury; acute and chronic pain; epilepsy and

seizures; olivopontocerebellar dementia; neuronal cell death; hypoxia-related neurodegeneration;

acute hypoxia; glutamate toxicity including glutamate neurotoxicity; cerebral ischemia; dementia

linked to meningitis and/or neurosis; cerebrovascular dementia; or dementia in an HIV-infected

patient.

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51. (Currently Amended) A method as claimed in claim 48 or 49, wherein the

neurodegenerative disorder is a peripheral neuropathy, including mononeuropathy, multiple

mononeuropathy or polyneuropathy, such as may be found in diabetes mellitus, Lyme disease or

uremia; peripheral neuropathy caused by a toxic agent; demyelinating disease such as acute or

chronic inflammatory polyneuropathy, leukodystrophies or Guillain-Barré syndrome; multiple

mononeuropathy secondary to a collagen vascular disorder; multiple mononeuropathy secondary

to sarcoidosis; multiple mononeuropathy secondary to a metabolic disease, or multiple

mononeuropathy secondary to an infectious disease.

52. (Currently Amended) A method as claimed in claim 46, 47 or 48, wherein the disorder is

inflammatory bowel disorder; bronchitis; asthma; acute pancreatitis; chronic pancreatitis;

allergies of various types; Alzheimer's disease; autoimmune disease such as rheumatoid arthritis,

systemic lupus erythematosus, glumerulonephritis, scleroderma, chronic thyroiditis, Graves's

disease, autoimmune gastritis, diabetes, autoimmune haemolytis anaemia, autoimmune

neutropaenia, thrombocytopenia, atopic dermatitis, chronic active hepatitis, myasthenia gravis,

multiple sclerosis, ulcerative colitis, Crohn's disease, psoriasis or graft vs host disease.

53. (Currently Amended) A method as claimed in any of claims 46-52, claim 46, wherein

one or more other active agent is administered to the individual simultaneously, subsequently or

sequentially to administering the compound.

54. (Original) A method as claimed in claim 53, wherein the other active agent is an anti-

inflammatory agent.

55. (Canceled)

56. (Canceled)

57. (Canceled)

58. (Canceled)

59. (Canceled)

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- 60. (Canceled)
- 61. (Canceled)
- 62. (Canceled)
- 63. (Currently Amended) An assay for determining the activity of the compounds as defined in any of claims 1-14, claim 1, comprising providing a system for assaying the activity and assaying the activity of a compound as defined in any of claims 1-14 claim 1.
- 64. (Original) An assay as claimed in claim 63, wherein the assay is for the JNK inhibiting activity of the compound, preferably for the JNK3-specific inhibiting activity of the compound.
- 65. (Currently Amended) An assay as claimed in claim 63 or 64, wherein the assay is a Scintillation Proximity Assay (SPA) using radiolabelled ATP, or is an ELISA.
- 66. (Currently Amended) A method of inhibiting the activity or function of a JNK, particularly JNK3, which method comprises exposing a JNK to a compound as defined in any of claims 1-14 or a composition as defined in any of claims 34-45 claim 1.
- 67. (Original) A method as claimed in claim 66, which is performed in a research model.
- 68. (Original) A method as claimed in claim 67, wherein the research model is an animal model.